

## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A method of treating an atherosclerotic target region of a coronary vessel in a patient, comprising  
delivering to the patient a photoatherolytic compound to cause accumulation of the compound in the target region,  
accessing the target region intraluminally with a guidewire,  
advancing over the guidewire a catheter having (i) a proximal main-body sleeve, (ii) a flexible, non-inflatable, translucent distal-end sleeve joined to the main-body sleeve at a catheter juncture, and (iii) an inner lumen extending through the two sleeves, said advancing being effective to position to the catheter's distal-sleeve within the target region,  
removing the guidewire from the catheter,  
introducing through the catheter a fiber-optic bundle having a light-diffusing tip, until said tip is positioned adjacent the catheter juncture,  
wherein the fiber-optic bundle is slidably associated with the catheter lumen;  
injecting a light-transmissive fluid through the catheter into the catheter's distal-end sleeve, ~~and~~  
irradiating the atherosclerotic vessel region by passing a laser light beam through the fiber optic bundle;  
wherein said beam is distributed along the catheter's distal-end sleeve, for transmission through the sleeve, by light scattering produced by (i) the light-diffusing tip, (ii) the light-transmissive fluid in the catheter's distal-end sleeve and (iii) the distal sleeve, and the scattered light transmitted through the sleeve is effective to photoactivate the photoatherolytic compound contained in the target region, ~~and wherein the proximal main-body sleeve of the catheter is associated with a handle that permits for intermittent injections of both a light transmissive fluid and a contrast fluid; and~~  
intermittently injecting a light transmissive fluid and a contrast fluid through the catheter into the catheter's distal-end sleeve.
2. (original) The method of claim 1, wherein the photoatherolytic compound is selected from the group consisting of a phycocyanin, a phthalocyanine, pheophorbide

derivative PH-1126, mono-L-aspartyl chlorin e6 (NPe6), hematoporphyrin derivative (HpD), benzoporphyrin derivative (BPD), Photofrin and Photofrin 2, protoporphyrin IX, and dihematoporphyrin-ester and -ether (DHE).

3. (original) The method of claim 1, wherein said injecting includes forcing a transparent or translucent aqueous solution through the catheter lumen.

4. (original) The method of claim 1, wherein said irradiating is carried out for a total of between about 10-20 minutes.

5. (original) The method of claim 4, wherein the catheter has a catheter wall port distal to said juncture, said port is positioned downstream of the target region in the coronary vessel, when the catheter is fully advanced, and said method further includes, at one or more intervals, during said irradiating step, retracting the catheter to position said port upstream the target, and thereby allow blood in said vessel to flow into and through said distal end region, to promote blood flow through the target region of the vessel at intervals during the treatment procedure.

6. (previously presented) Apparatus for use in treating an atherosclerotic target region of a coronary vessel in a patient, comprising

a guidewire for accessing the target region intraluminally,

a catheter having (i) a proximal main-body sleeve associated with a handle, (ii) a flexible, non-inflatable, translucent distal-end sleeve joined to the main-body sleeve at a catheter juncture, and (iii) an inner lumen extending through the two sleeves, through which lumen the catheter can be advanced over the guidewire to position the catheter's distal-end sleeve within the target region,

a fiber-optic bundle having a light-diffusing tip, said bundle being adapted for introduction into and slidably associated with the catheter lumen after the catheter's distal-end sleeve is positioned within the target region, and the guidewire is removed;

a proximal-end catheter port through which a light-transmissive fluid can be injected through the catheter into the catheter's distal-end sleeve, and

a proximal-end optical connector to which the fiber-optic bundle can be connected to a laser source, for irradiating the atherosclerotic target region by passing a laser light beam through the fiber optic bundle,

such that the laser beam is distributed along the catheter's distal-end sleeve, for transmission through the sleeve, by light scattering produced by (i) the light-diffusing tip, (ii) the light-transmissive fluid injected into the catheter's distal-end sleeve and (iii) the distal-end sleeve, and where the scattered light transmitted through the sleeve is effective to photoactivate a photoatherolytic compound contained in the target region; and  
wherein the handle permits for intermittent injections of both a light-transmissive fluid and a contrast fluid.

7. (original) The apparatus of claim 6, wherein said catheter has an inner-lumen diameter of between about .45 and .6 mm.

8. (original) The apparatus of claim 6, wherein the optic fiber bundle is formed of a plurality of light fibers encased in an outer sleeve for relative axial fiber sliding movement, to enhance the flexibility of the fiber bundle.

9. (original) The apparatus of claim 6, wherein said catheter has a wall port downstream of said juncture, and located to allow blood in the patient's vessel to flow into and through said distal end sleeve, with the catheter distal-end sleeve placed in the target region, and withdrawn to place the port just upstream of the target region.

10. (original) Apparatus for use in treating an atherosclerotic target region of a coronary vessel in a patient, comprising  
a guidewire for accessing the target region intraluminally,  
a catheter having (i) a proximal main-body sleeve associated with a handle, (ii) a flexible, non-inflatable, translucent distal-end sleeve joined to the main-body sleeve at a catheter juncture, and (iii) an inner lumen extending through the two sleeves, through which lumen the catheter can be advanced over the guidewire to position the catheter's distal-end sleeve within the target region,  
a fiber-optic bundle having a light-diffusing tip, said bundle being adapted for introduction into and slidably associated with the catheter lumen after the catheter's distal-end sleeve is positioned within the target region, and the guidewire is removed;  
a proximal-end catheter port through which a light-transmissive fluid can be injected through the catheter into the catheter's distal-end sleeve, and

a proximal-end optical connector to which the fiber-optic bundle can be connected to a laser source, for irradiating the atherosclerotic target region by passing a laser light beam through the fiber optic bundle,

such that the laser beam is distributed along the catheter's distal-end sleeve, for transmission through the sleeve, by light scattering produced by (i) the light-diffusing tip, (ii) the light-transmissive fluid injected into the catheter's distal-end sleeve and (iii) the distal-end sleeve, and where the scattered light transmitted through the sleeve is effective to photoactivate a photoatherolytic compound contained in the target region; and

wherein the handle comprises a first syringe containing a light-transmissive fluid and a second syringe containing a contrast fluid.

11. (original) The apparatus of claim 10, wherein the handle permits for intermittent injections of both the light-transmissive fluid and the contrast fluid.

12. (original) The apparatus of claim 11, wherein the handle further comprises a timer and a switching device configured to automatically inject the light-transmissive fluid and contrast fluid.

13. (original) The apparatus of claim 10, wherein the handle further comprises first and second indicator lights configured to indicate the flow of the light-transmissive fluid and the contrast fluid.

14. (currently amended) ~~The apparatus of claim 13,~~ Apparatus for use in treating an atherosclerotic target region of a coronary vessel in a patient, comprising  
a guidewire for accessing the target region intraluminally,  
a catheter having (i) a proximal main-body sleeve associated with a handle, (ii) a flexible, non-inflatable, translucent distal-end sleeve joined to the main-body sleeve at a catheter juncture, and (iii) an inner lumen extending through the two sleeves, through which lumen the catheter can be advanced over the guidewire to position the catheter's distal-end sleeve within the target region,  
a fiber-optic bundle having a light-diffusing tip, said bundle being adapted for introduction into and slidably associated with the catheter lumen after the catheter's distal-end sleeve is positioned within the target region, and the guidewire is removed;

a proximal-end catheter port through which a light-transmissive fluid can be injected through the catheter into the catheter's distal-end sleeve, and

a proximal-end optical connector to which the fiber-optic bundle can be connected to a laser source, for irradiating the atherosclerotic target region by passing a laser light beam through the fiber optic bundle,

such that the laser beam is distributed along the catheter's distal-end sleeve, for transmission through the sleeve, by light scattering produced by (i) the light-diffusing tip, (ii) the light-transmissive fluid injected into the catheter's distal-end sleeve and (iii) the distal-end sleeve, and where the scattered light transmitted through the sleeve is effective to photoactivate a photoatherolytic compound contained in the target region; and

wherein the handle comprises a first syringe containing a light-transmissive fluid and a second syringe containing a contrast fluid;

wherein the handle further comprises first and second indicator lights configured to indicate the flow of the light-transmissive fluid and the contrast fluid;

wherein the catheter further comprises a sensor, and

wherein the sensor is configured to activate the first and second indicator lights.

15. (original) The apparatus of claim 14, wherein the sensor is located at the distal end of the catheter.